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(54) **Pharmaceutical formulations of cefaclor**

Pharmazeutische Zubereitungen enthaltend Cefaclor

Formulations pharmaceutiques contenant du cefaclore

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Description

FIELD OF THE INVENTION

5 [0001] This invention refers to pharmaceutical formulations containing cefaclor suitable for the manufacture of solid pharmaceutical forms for oral administration. In particular, the invention refers to pharmaceutical formulations containing cefaclor.

BACKGROUND OF THE INVENTION

10 [0002] Cefaclor or 3-chloro-7-D-(phenylglycinamide)-3-cephem-4-carboxylic acid is a semisynthetic cephalosporinic antibiotic described for example in US Patent no. 3,925,372 and German Patent No. DE 2,408,698 (Eli Lilly & Co.). Its bactericide action is based on its capacity to inhibit cell wall synthesis. Cefaclor is indicated for treatment of infections caused by sensitive strains of numerous organisms, particularly *Streptococcus* and *Staphylococcus*.

15 [0003] Pharmaceutical forms currently available for administration of cefaclor include capsules, retard tablets and suspensions, both in phial and sachet form.

[0004] There are a number of limitations and drawbacks to the use of capsules affecting the following:

- their administration to the patient, which may be limited because some may have problems in swallowing them, particularly children and the elderly who may even be unable to do so; and
- the fact that they permit only a single dosage.

[0005] On the other hand, the drawback to cefaclor administration in suspension (sachets) is that, due to its saccharose content, its potential use is limited in diabetic patients, who must take the appropriate precautions.

25 [0006] Moreover, added to this drawback caused by the saccharose content, administration of cefaclor in suspension (phial) has further disadvantages which may be summarised as follows:

- risk of accidental overdose due to uncontrolled consumption, particularly in children; and
- the difficulties of handling and transport because of the volume involved,

30 leading to some risk of failure to complete the therapy, with the attendant loss of efficacy.

[0007] Retard tablets have the drawback of not permitting double dosage of the product; moreover, they are not suitable for patients with difficulties in ingesting solid forms.

35 [0008] Existing forms of administration of cefaclor do not in general meet entirely satisfactorily some requirements considered desirable in the treatment of bacterial infections, for example their application to any patient in conditions such as to ensure that the therapy is completed, thereby enhancing its efficacy.

[0009] There is thus a requirement for new pharmaceutical forms for administering cefaclor which solve these problems, making it easier for the patient to administer them, and so that they can be used with diabetic patients without causing additional difficulties, so increasing the efficacy of treatment. This invention provides a solution to these problems, by furnishing new pharmaceutical formulations containing cefaclor which are suitable for the manufacture of dispersible tablets.

40 [0010] Dispersible tablets are solid pharmaceutical forms for oral administration which must disintegrate in less than three minutes in water at 19°C - 21°C and disperse evenly in the water. The dispersion uniformity trial involves placing two tablets in 100 ml of water and shaking them until they are completely disintegrated; they must disperse so as to pass through a screen with a nominal mesh size of 710 microns (*British Pharmacopea* Vol. II, 1988).

45 [0011] Dispersible tablets are known which contain antibiotics belonging to the synthetic penicillins group (amoxycillin) and anti-inflammatories (pyroxycam), but none is known which contains an antibiotic belonging to the synthetic cephalosporin group, such as cefaclor.

50 [0012] The subject of this invention is new pharmaceutical formulations containing cefaclor suitable for the manufacture of dispersible tablets.

A DETAILED DESCRIPTION OF THE INVENTION

55 [0013] The preparation of formulations suitable for manufacturing dispersible tablets requires both study of the physical-chemical incompatibilities of the active ingredient and a search for the right excipients making it possible to comply with the requirements of the various Pharmacopeas. Account must also be taken of the procedure to be used for the manufacture of such dispersible tablets, since the excipients and coadjuvants of the formulation will in large part depend on the process selected for the manufacture of such dispersible tablets. For reasons to be mentioned subsequently,

direct compression was the process selected for the manufacture of the tablets.

[0014] The parameters defining dispersible tablets are as follows:

- i) Their high rate of disintegration in water, and
- ii) The uniform distribution of the particles into which they disintegrate.

[0015] Disintegration rate and uniformity of dispersion are also dependent on both the coadjuvants and the active ingredient. Thus disintegration, as a measure of the release of the active ingredient in compressed pharmaceutical preparations, is the critical parameter for the development of dispersible forms. Thus the selection of coadjuvants in the preparation of dispersible tablets is the most important phase of the Galenic research. The properties and the quality of the finished tablet depend in large part on the coadjuvants it incorporates so that the correct choice of coadjuvant is of the greatest importance, as is the manufacturing process since the type of coadjuvant may be selected depending on the technique used.

[0016] New pharmaceutical formulations of cefaclor, suitable for manufacturing dispersible tablets provided by this invention, take account of these considerations and, in addition to the active ingredient, contain adequate amounts of disintegrator, diluents, lubricants, antiadherents, sweeteners, fragrances and, optionally, flavorings, flattening agents and colorings. In addition, in a particular implementation of this invention, new pharmaceutical formulations of cefaclor are provided which incorporate an effervescent pair.

[0017] Cefaclor is the active ingredient of the formulations in this invention. As used in this description, the term "cefaclor" is intended to include not only the free acid form but also its hydrates and pharmaceutically acceptable salts. The Cefaclor may be present in the formulation of an amount between 35% and 50% by weight of the total formulation weight. The cefaclor can be prepared as described for example in US Patent no. 3,925,372 and German Patent No. DE 2,408,698 (Eli Lilly & Co.).

[0018] Because the critical parameter of dispersible tablets is their rate of disintegration in water, the choice of the right disintegrator is one of the most important stages. As used in this description, the term "disintegrator" refers to an agent which produces a surface increase so that the active ingredient of the tablet is released very quickly. Glycolate sodium starch, alone or together with carboxymethylcellulose, polymeric derivatives of acrylic acid and, preferably, croscopovidone are suitable disintegrators for the formulations in this invention.

[0019] Glycolate sodium starch can be used in proportions of 5% and more by weight of the total formulation weight and, for preference, at concentrations between 10% and 21%. In addition, mixtures of glycolate sodium starch and sodium carboxymethylcellulose may be used in amounts of approximately 14% by weight of glycolate sodium starch and approximately 10% of sodium carboxymethylcellulose, in both cases in relation to the total weight of the formulation.

[0020] The polymeric derivative of acrylic acid may be of medium or high viscosity, preferably high, and may be used in a proportion of approximately 10% by weight of the total formulation weight.

[0021] The preferred disintegrator is a croscopovidone (insoluble polyvinylpyrrolidone [PVP] obtained by polymerization of vinylpyrrolidone). This polymer can be included in the formulation in a proportion of approximately 10% by weight of the total formulation weight. It is believed that the high disintegrating action of the reticulated and insoluble PVP is due to its hydration capacity (water adsorption) which means that a very rapid tablet disintegration rate is attained with the resulting improved dissolution of the cefaclor in water.

[0022] On the other hand, the selection of the direct compression technique to manufacture dispersible tablets involves a further advantage in the choice of excipients. The possibility of using the disintegrator in extragranular form enhances its swelling effect since the disintegrating effect is not altered either by wetting or by drying.

[0023] In the sense used in this description, the term "diluents" includes excipients which facilitate the compression of powdery materials and give the tablets strength. Microcrystalline cellulose and dry flowing starch and mixtures thereof are suitable diluents.

[0024] The following are examples of suitable diluents for the formulations in this invention:

- 1) Microcrystalline cellulose, which provides the powder mixture with highly appropriate fluidity and compressibility characteristics. This diluent makes it possible to manufacture tablets of a high level of purity, using the direct compression technique. It also acts as a binder, to give strong tablets of suitable hardness, while its absorption capacity contributes to short disintegration times. Of the different types of microcrystalline cellulose available on the market, AVICEL PH102 (mean particle size 90 microns) is preferred; while the others have similar characteristics in terms of their capacity to facilitate direct compression, AVICEL PH102 makes direct compression more straightforward thanks to the fluidity it confers on the mixture and, because of its particle size, it facilitates direct compression of fine powder mixtures (as in the formulations of this invention). The microcrystalline cellulose may be present in the formulation at between 24% and 46% by weight of total formulation weight, and
- 2) dry flowing starch, due to its diluent and binding capacity in direct compression. It may be present in the formulation in a quantity of approximately 39% by weight of total formulation weight. However, tablets made with this

diluent are not very hard, and this affects their friability negatively: for this reason, tablets containing microcrystalline cellulose as diluent are preferred.

[0025] Because of the very high percentage of microcrystalline cellulose (between 24% and 46% by weight of total formulation weight) tablets can be obtained with weights of between 1140 and 1150 mg, with high cellulose percentages, of the order of 38% to 46%, while with somewhat lower percentages - of the order of 35%-36% - tablets were obtained with a final weight of the order of 1125 mg each. Finally, when the percentage of microcrystalline cellulose is approximately 28%, tablets can be obtained with final weights of the order of 1130 - 1140 mg.

[0026] The term "lubricant" as used in this description includes excipients which reduce interparticle friction inside the tablet, reducing the reaction forces appearing on the walls of the matrix. Stearyl sodium fumarate, a hydrophylic lubricant, may be used for preference as lubricant suitable for the formulations in this invention. This coadjuvant can be added to the formulations in this invention at a rate of less than 2% by weight in relation to the total weight of the formulation and, for preference, between 0.4% and 1.5% by weight of the total formulation weight.

[0027] The inclusion of this excipient enhances the slipping of the formulation to be compressed. It also ensures even filling of the space in the matrix so that there is very little tablet weight variation.

[0028] Standard stearic acid salts are not suitable since, for example, magnesium stearate does not adsorb water, giving a solution of most unpleasant appearance, with the formation of a "halo" on the surface.

[0029] The term "antiadherent" as used in this description includes excipients which prevent particle adhesion, so avoiding or reducing compacting and limiting friction between them. Colloidal silicon dioxide can be used as a suitable antiadherent for the formulations in this invention: because of its large specific surface, this raw material is a very good regulator of powder flow and also acts as adsorbent, capturing the humidity which would be taken up by the cefaclor, so slowing the degradation of the active ingredient by hydrolysis.

[0030] This coadjuvant can be incorporated at a percentage of less than 5% by weight of the total formulation weight and, for preference, between 0.2% and 1.5%.

[0031] The formulations in this invention may also contain sweeteners, fragrances and flavorings. Sodium saccharin may be used as artificial sweetener at less than 1% by weight of total weight of the formulation and, for preference, between 0.1% and 0.4% by weight, or aspartame at less than 1% by weight of total formulation weight and, for preference, between 0.2% and 0.75% by weight in relation to total weight of the formulation.

[0032] Strawberry fragrance may be used for preference as fragrance, for example that identified as 52,312 AP05.15 Firmenich, at between 3% and 6% by weight of the total formulation weight.

[0033] Anhydrous citric acid can be used for preference as flavoring, at between 2% and 4% by weight of total formulation weight.

[0034] Additionally and optionally, the formulations in this invention may contain a flattening agent and a colorant or combination of colorants to enhance the physical appearance of the solution to be obtained and to give it a uniform color. Titanium dioxide (E-171) may be used as an opacifier, in an amount less than 2% by weight of the total formulation weight, and for preference approximately 1.5% by weight of total formulation weight. Its use is not however essential.

[0035] Individual colorants or combinations may be used, preferably to obtain a red-pink solution which can be associated with the fragrance to be used in the tablets (strawberry) and which also provide the final suspension with a pleasant appearance. These objectives may be attained by including less than 1% by weight of the Red F, D and C No. 3 - erythrosine (E-127) coloring [Merck Index, 11th Edition, 1989, Rahway, N.J., USA].

[0036] On the other hand, in a particular and alternative implementation of this invention, new pharmaceutical formulations of cefaclor are provided which also incorporate a pair of compounds which produce an effervescent effect. By including the effervescent pair, the tablet's disintegration rate can be increased. In general, an effervescent pair consists of an effervescent base, such as an alkaline or earth alkaline metal carbonate or bicarbonate and an acid which, on reacting with the effervescent base, produces carbon dioxide. In the new cefaclor formulations in this invention, any of the effervescent pairs normally used to make effervescent tablets can be employed, preferably that consisting of citric acid and calcium carbonate.

[0037] The cefaclor formulations obtained from this invention may be prepared easily by screening the appropriate amounts of the different excipients and coadjuvants and placing them in a suitable mixer. The active ingredient is then added and mixed until homogeneous, to give a free-flowing powder.

[0038] These new formulations can be used to make dispersible tablets containing cefaclor as active ingredient.

[0039] As already stated, the manufacturing process for the tablet plays a very important role in the design of the pharmaceutical formulation. The formation of the tablet casing may be based on a granulate (an agglomerated material made of powder particles to which a binder is added), or on a previously untreated powder mixture (direct compression). Coadjuvants are selected according to the technique selected. Because dispersible tablets are very sensitive to damp and their stability is compromised by granulation operations, direct compression is the preferred technique, being the one which offers most advantages: on the one hand, manufacture is rapid, depending neither on granulation or drying and, on the other, it avoids possible degradation (due to hydrolysis) of the active ingredient during granulation. Risk of

contamination is also reduced. However, perhaps the most significant advantage is that directly compressed tablets normally disintegrate more rapidly than those made by wet granulation which require the addition of binding agents that slow the disintegration rate.

[0040] While direct compression may cause some drawbacks, such as problems of uniformity of mixture and dosage, fluidity and compressibility, surprisingly, with the formulations in this invention, none of these problems arose. In fact, the tablets varied very little in weight and content of active ingredient. Compressibility was acceptable and tablet hardness was within the required limits.

[0041] Dispersible tablets containing cefaclor may be manufactured by standard processes, for example in a conventional rotary or eccentric compressing machine which compresses the prepared and screened pharmaceutical formulation fed to the machine.

[0042] Dispersible tablets containing cefaclor provided by this invention are solid, intended for oral use, of uniform appearance, and with sufficient mechanical strength to withstand possible damage from storage and transport. The active ingredient is distributed evenly in the pharmaceutical form and the disintegration rate in water is high (within three minutes in water at 19°C - 21°C). Likewise, the level of disintegration (or fineness of the particles in to which the product disintegrates) is suitable, and in line with the requirements of the various *Pharmacopeas*.

[0043] The use of dispersible tablets containing cefaclor provides a series of benefits over known and habitual forms of administration of this active ingredient, including the following:

- They are suitable for treating patients with difficulties in ingesting solid forms;
- They may be used by diabetic patients since they do not contain saccharose;
- Dosage is flexible and reasonably accurate following dissolution in the volume of water desired by the patient.
- The resulting solutions are of suitable organoleptic characteristics, acceptable to patients.
- Their shape, size and reduced volume allow them to be presented in blister form, which is a benefit to the patient, enhancing ease of handling and carrying, to ensure that the patient completes the therapy, thereby raising the efficacy of treatment; and
- There is less risk of accidental intoxication due to overdose, making them less hazardous, particularly to children.

[0044] The following Examples illustrate specific implementations of the invention and should not be construed as limiting it. Said Examples use dry flowing starch [GORMASO], crospovidone (KOLLIDON CL®) [BASF] and microcrystalline cellulose (AVICEL PH 102®) [FMC FORET]. Both high- and medium-viscosity acrylic derivatives consist of a copolymer of methacrylic acid and methyl methacrylate in a ratio of approximately 7:3, made by ROHM PHARMA. The difference between the two types of acrylic derivatives is due to the different viscosity of the gel they form.

EXAMPLE 1

[0045] Dispersible tablets were prepared from the following pharmaceutical formulation:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	37.35
Glycolate sodium starch	70.00	5.00
CMC*, AVICEL PH102	612.99	43.78
Stearyl sodium fumarate	14.00	1.00
Colloidal silicon dioxide	70.00	5.00
Aspartame	10.00	0.71
Strawberry fragrance	50.00	3.58
Anhydrous citric acid	50.00	3.58

*CMC Microcrystalline cellulose

[0046] The process begins with the weighing of all the raw materials separately then screening them as a security measure. After screening, the excipients are placed in a suitable mixer, the cefaclor is then added and re-mixed until homogenous.

[0047] The mixed powders are passed several times through a 0.7 mm mesh screen. Compression follows, with periodic controls during the process, noting the results obtained on the associated control cards. The powder flows satisfactorily and compresses without difficulties. At the end of the process, representative samples are taken for analysis (from the beginning, middle and end of the batch) using a statistical sampling procedure.

[0048] Dispersible tablets were obtained with the following characteristics:

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Individual tablet weight: 1400 mg \pm 5%

Weight of 10 tablets: 14 g \pm 3%

Hardness: 7 Kg

Height: approximately 5.6 mm

Diameter: 17.15 mm

Disintegration in water at 19°C - 21°C: <3 min

Appearance: white with some transparent surface specks

Flavor: pleasant (quite neutral)

Suspension: once disintegrated and shaken, remains in suspension approximately 2 minutes.

EXAMPLE 2

[0049] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	41.84
Glycolate sodium starch	62.50	5.00
CMC*, AVICEL PH102	544.49	43.56
Stearyl sodium fumarate	12.50	1.00
Colloidal silicon dioxide	62.50	5.00
Aspartame	5.00	0.40
Strawberry fragrance	40.00	3.20

*CMC Microcrystalline cellulose

Dispersible tablets were obtained with the following characteristics:

[0050]

Individual tablet weight: 1250 mg \pm 5%

Weight of 10 tablets: 12.5 g \pm 3%

Hardness: 7.8 Kgf

Height: approximately 5.5 mm

Diameter: 17.15 mm

Disintegration in water at 19°C - 21°C: <3 min Flavor: pleasant (quite neutral)

Suspension: once disintegrated and shaken, remains in suspension approximately 2 minutes.

EXAMPLE 3

[0051] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	41.84
Glycolate sodium starch	62.50	5.00
CMC*, AVICEL PH102	574.49	45.96
Stearyl sodium fumarate	12.50	1.00
Colloidal silicon dioxide	12.50	1.00
Sodium saccharin	5.00	0.40
Strawberry fragrance	60.00	4.80

*CMC Microcrystalline cellulose

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Dispersible tablets were obtained with the following characteristics:

[0052]

- 5 Individual tablet weight: 1250 mg \pm 5%
Weight of 10 tablets: 12.5 g \pm 3%
Hardness: 8 Kgf
Height: approximately 5.2 mm
Diameter: 17.15 mm
10 Disintegration in water at 19°C - 21°C: <3 min
Appearance: white-ish
Flavor: pleasant
Suspension: once disintegrated and shaken, remains in suspension approximately 2 minutes.

15 EXAMPLE 4

[0053] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.00	41.84
Glycolate sodium starch	62.50	5.00
CMC*, AVICEL PH102	562.00	44.96
25 Stearyl sodium fumarate	12.50	1.00
Colloidal silicon dioxide	12.50	1.00
Sodium saccharin	5.00	0.40
Strawberry fragrance	60.00	4.80
30 Titanium dioxide	12.50	1.00

*CMC Microcrystalline cellulose

Dispersible tablets were obtained with the following characteristics:

35 [0054]

- Individual tablet weight: 1250 mg \pm 5%
Weight of 10 tablets: 12.5 g \pm 3%
Hardness: 10 Kgf
40 Height: approximately 5.1 mm
Diameter: 17.15 mm
Disintegration in water at 19°C - 21°C: <3 min
Appearance: white-ish
Flavor: pleasant
45 Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 5

- 50 [0055] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	45.49
55 Glycolate sodium starch	115.00	10.00
Dry flowing starch	445.00	38.70
Stearyl sodium fumarate	6.25	0.54

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(continued)

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Colloidal silicon dioxide	6.25	0.54
Sodium saccharin	3.00	0.27
Strawberry fragrance	45.00	3.92
Titanium dioxide	6.25	0.54

[0056] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1149.76 mg \pm 5%

Weight of 10 tablets: 11.5 g \pm 3%

Hardness: 8 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): 0.9%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 6

[0057] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	48.29
Glycolate sodium starch	163.00	15.05
CMC*, AVICEL PH102	345.00	31.85
Stearyl sodium fumarate	5.60	0.52
Colloidal silicon dioxide	2.20	0.20
Sodium saccharin	3.00	0.28
Strawberry fragrance	38.00	3.51
Titanium dioxide	3.30	0.30

*CMC Microcrystalline cellulose

[0058] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1083.11 mg \pm 5%

Weight of 10 tablets: 10.8 g \pm 3%

Hardness: 8 - 9 Kgp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): 0.8%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 7

[0059] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

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COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	45.88
Glycolate sodium starch	163.00	14.30
Sodium carboxymethylcellulose	120.00	10.53
CMC*, AVICEL PH102	275.00	24.13
Stearyl sodium fumarate	5.60	0.49
Colloidal silicon dioxide	6.00	0.53
Sodium saccharin	2.00	0.17
Strawberry fragrance	42.00	3.68
Titanium dioxide	3.30	0.29

*CMC Microcrystalline cellulose

[0060] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1139.91 mg \pm 5%

Weight of 10 tablets: 11.4 g \pm 3%

Hardness: 8 - 9 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): 0.8%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 8

[0061] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	45.88
Glycolate sodium starch	110.01	9.65
Calcium carbonate	14.02	1.23
Anhydrous citric acid	40.01	3.51
CMC*, AVICEL PH102	390.02	34.21
Stearyl sodium fumarate	5.59	0.50
Colloidal silicon dioxide	11.97	1.05
Sodium saccharin	2.05	0.18
Strawberry fragrance	40.01	3.50
Titanium dioxide	3.31	0.29

*CMC Microcrystalline cellulose

[0062] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1140 mg \pm 5%

Weight of 10 tablets: 11.4 g \pm 3%

Hardness: 9 - 10 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): 0.6%

Appearance: white-ish

Flavor: pleasant

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EXAMPLE 9

[0063] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.10
Glycolate sodium starch	171.00	15.07
CMC*, AVICEL PH102	345.00	31.35
Stearyl sodium fumarate	17.10	1.50
Colloidal silicon dioxide	17.10	1.50
Sodium saccharin	2.28	0.21
Strawberry fragrance	39.90	3.52
Titanium dioxide	8.55	0.75

*CMC Microcrystalline cellulose

[0064] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1134.62 mg \pm 5%

Weight of 10 tablets: 11.3 g \pm 3%

Hardness: 12.8 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 10

[0065] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.10
Glycolate sodium starch	171.00	15.07
CMC*, AVICEL PH102	355.68	31.35
Stearyl sodium fumarate	17.10	1.50
Colloidal silicon dioxide	17.10	1.50
Aspartame	2.28	0.21
Strawberry fragrance	39.90	3.52
Titanium dioxide	8.55	0.75

*CMC Microcrystalline cellulose

[0066] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1134.62 mg \pm 5%

Weight of 10 tablets: 11.3 g \pm 3%

Hardness: 12.8 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

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minutes).

EXAMPLE 11

- 5 [0067] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	45.96
Glycolate sodium starch	205.20	18.03
CMC*, AVICEL PH102	324.90	28.55
Stearyl sodium fumarate	17.10	1.50
Colloidal silicon dioxide	17.10	1.50
Sodium saccharin	2.28	0.20
Strawberry fragrance	39.90	3.51
Titanium dioxide	8.55	0.75

*CMC Microcrystalline cellulose

- 20 [0068] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1138.04 mg \pm 5%

Weight of 10 tablets: 11.4 g \pm 3%

Hardness: 13.0 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 12

- 35 [0069] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	45.96
Glycolate sodium starch	228.00	20.03
CMC*, AVICEL PH102	302.10	26.55
Stearyl sodium fumarate	17.10	1.50
Colloidal silicon dioxide	17.10	1.50
Sodium saccharin	2.28	0.20
Strawberry fragrance	39.90	3.51
Titanium dioxide	8.55	0.75

*CMC Microcrystalline cellulose

- 50 [0070] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1138.04 mg \pm 5%

Weight of 10 tablets: 11.4 g \pm 3%

Hardness: 10.7 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: white-ish

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Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 13

[0071] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.66
Acrylic derivate (medium viscosity)	114.00	10.17
CMC*, AVICEL PH102	399.00	35.60
Stearyl sodium fumarate	17.10	1.50
Colloidal silicon dioxide	17.10	1.50
Sodium saccharin	2.28	0.21
Strawberry fragrance	39.90	3.56
Titanium dioxide	8.55	0.76

*CMC Microcrystalline cellulose

[0072] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1120.94 mg \pm 5%

Weight of 10 tablets: 11.2 g \pm 3%

Hardness: 12.27 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 14

[0073] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.26
Acrylic derivate (high viscosity)	114.00	10.08
CMC*, AVICEL PH102	399.00	35.30
Stearyl sodium fumarate	17.10	1.51
Colloidal silicon dioxide	17.10	1.51
Sodium saccharin	3.42	0.30
Strawberry fragrance	39.90	3.53
Titanium dioxide	17.10	1.51

*CMC Microcrystalline cellulose

[0074] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1130.63 mg \pm 5%

Weight of 10 tablets: 11.3 g \pm 3%

Hardness: 13.65 Kp

Disintegration in water at 19°C - 21°C: <3 min

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Appearance: white-ish

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 15

[0075] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.11
Acrylic derivate (high viscosity)	114.00	10.05
CMC*, AVICEL PH102	399.00	35.17
Stearyl sodium fumarate	17.10	1.51
Colloidal silicon dioxide	17.10	1.51
Sodium saccharin	4.10	0.36
Strawberry fragrance	39.90	3.52
Titanium dioxide	17.10	1.51
Erythrosine	3.00	0.26

*CMC Microcrystalline cellulose

[0076] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1134.31 mg \pm 5%

Weight of 10 tablets: 11.3 g \pm 3%

Hardness: 13 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: pale rose color

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 16

[0077] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.24
Acrylic derivate (high viscosity)	114.00	10.08
CMC*, AVICEL PH102	400.00	35.07
Stearyl sodium fumarate	17.10	1.51
Colloidal silicon dioxide	17.10	1.51
Sodium saccharin	4.10	0.36
Strawberry fragrance	40.00	3.53
Yellow F, D and C No. 6	4.00	0.35
Erythrosine	4.00	0.35

*CMC Microcrystalline cellulose

[0078] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1131.31 mg \pm 5%

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Weight of 10 tablets: 11.3 g \pm 3%

Hardness: 15.74 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: pale rose color

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 17

[0079] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.48
Acrylic derivate (high viscosity)	114.00	10.13
CMC*, AVICEL PH102	400.00	35.34
Stearyl sodium fumarate	17.10	1.52
Colloidal silicon dioxide	17.10	1.52
Sodium saccharin	4.10	0.36
Strawberry fragrance	40.00	3.56
Erythrosine	10.00	0.89

*CMC Microcrystalline cellulose

[0080] Dispersible tablets were obtained with the following characteristics:

Individual tablet weight: 1125.31 mg \pm 5%

Weight of 10 tablets: 11.2 g \pm 3%

Hardness: 15 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: pale rose color

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

EXAMPLE 18

[0081] Dispersible tablets were prepared from the following pharmaceutical formulation, using the procedure described in Example 1:

COMPONENTS	WEIGHT (mg)	% BY WEIGHT
Cefaclor	523.01	46.48
Crospovidone	114.00	10.13
CMC*, AVICEL PH102	400.00	35.34
Stearyl sodium fumarate	17.10	1.52
Colloidal silicon dioxide	17.10	1.52
Sodium saccharin	4.10	0.36
Strawberry fragrance	40.00	3.56
Erythrosine	10.00	0.89

*CMC Microcrystalline cellulose

[0082] Dispersible tablets were obtained with the following characteristics:

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Individual tablet weight: 1125.31 mg \pm 5%

Weight of 10 tablets: 11.2 g \pm 3%

Hardness: 14.78 Kp

Disintegration in water at 19°C - 21°C: <3 min

Friability (10 tablets): <0.5%

Appearance: pale rose color

Flavor: pleasant

Suspension: once disintegrated and shaken, appears to remain in suspension for a long period (approximately 30 minutes).

Claims

1. A pharmaceutical formulation suitable for manufacturing dispersible tablets by direct compression, with cefaclor as active ingredient in a quantity between 35% and 50% by weight of total formulation weight, with suitable excipients and coadjuvants.
2. A formulation as set forth in claim 1 in which said suitable excipients and coadjuvants comprise disintegrators, diluents, lubricants, antiadherents, sweeteners, fragrances and, optionally, flavorings, opacifiers and colorants.
3. A formulation as set forth in claim 2 in which the disintegrator is glycolate sodium starch, acrylic derivates, mixtures of glycolate sodium starch and carboxymethylcellulose and, for preference, crospovidone.
4. A formulation as set forth in claim 3 comprising glycolate sodium starch in an amount between 5% and 21% by weight of total formulation weight.
5. A formulation as set forth in claim 3 comprising acrylic acid derivates in an amount of approximately 10% by weight of total formulation weight.
6. A formulation as set forth in claim 3 comprising crospovidone in an amount of approximately 10% by weight of total formulation weight.
7. A formulation as set forth in claim 3 comprising a mixture of (i) glycolate sodium starch in an amount of approximately 14% by weight of total formulation weight, and (ii) carboxymethylcellulose in an amount of approximately 10% by weight of total formulation weight, as disintegrator.
8. A formulation as set forth in claim 2 comprising microcrystalline cellulose, dry flowing starch and their mixtures, as diluent.
9. A formulation as set forth in claim 8 comprising microcrystalline cellulose in an amount between 24% and 46% by weight of total formulation weight.
10. A formulation as set forth in claim 9 in which said microcrystalline cellulose has a mean particle size of approximately 90 microns.
11. A formulation as set forth in claim 8 comprising dry flowing starch in an amount of approximately 39% by weight of total formulation weight.
12. A formulation as set forth in claim 2 comprising stearyl sodium fumarate as lubricant, in an amount of less than 2% by weight of total formulation weight.
13. A formulation as set forth in claim 2 comprising colloidal silicon dioxide as antiadherent, in an amount of less than 5% by weight of total formulation weight.
14. A formulation as set forth in claim 2 comprising aspartame, sodium saccharin or their mixtures as sweetener, in an amount of less than 1% by weight of total formulation weight.
15. A formulation as set forth in claim 14 comprising sodium saccharin in an amount between 0.1% and 0.4% by weight

of total formulation weight.

16. A formulation as set forth in claim 14 comprising aspartame in an amount between 0.2% and 0.7% by weight of total formulation weight.
17. A formulation as set forth in claim 2 comprising a strawberry fragrance, in an amount between 3% and 5% by weight of total formulation weight.
18. A formulation as set forth in claim 2 comprising anhydrous citric acid as flavoring, in an amount between 2% and 4% by weight of total formulation weight.
19. A formulation as set forth in claim 2 comprising titanium dioxide as opacifier, in an amount of less than 2% by weight of total formulation weight.
20. A formulation as set forth in claim 2 comprising Red F, D and C No. 3 - Erythrosine as colorant, in an amount of less than 1% by weight of total formulation weight.
21. A formulation as set forth in claim 1 which includes a pair of compounds producing an effervescent effect.
22. A formulation as set forth in claim 21 wherein said effervescent pair consists of citric acid and calcium carbonate.

Patentansprüche

1. Pharmazeutische Zubereitung, die für die Herstellung dispergierbarer Tabletten durch direkte Komprimierung geeignet ist, mit Cefaclor in einer Menge von zwischen 35 und 50 Gew.-% des Gesamtgewichtes der Zubereitung als aktivem Bestandteil und geeigneten Arzneimittelträgern und Coadjuvantien.
2. Zubereitung nach Anspruch 1, worin die geeigneten Arzneimittelträger und Coadjuvantien Desintegrationsmittel, Verdünnungsmittel, Gleitmittel, Antihafmittel, Süßungsmittel, Duftstoffe und wahlweise Aromastoffe, Trübungsmittel und Farbstoffe umfassen.
3. Zubereitung nach Anspruch 2, worin das Desintegrationsmittel Glykolatnatrium-Stärke, Acrylderivate, Mischungen von Glykolatnatrium-Stärke und Carboxymethylcellulose und bevorzugt Crospovidon ist.
4. Zubereitung nach Anspruch 3, die Glykolatnatrium-Stärke in einer Menge zwischen 5 und 21 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
5. Zubereitung nach Anspruch 3, die Acrylsäurederivate in einer Menge von ungefähr 10 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
6. Zubereitung nach Anspruch 3, die Crospovidon in einer Menge von ungefähr 10 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
7. Zubereitung nach Anspruch 3, die eine Mischung von (i) Glykolatnatrium-Stärke in einer Menge von ungefähr 14 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung und (ii) Carboxymethylcellulose in einer Menge von ungefähr 10 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung als Desintegrationsmittel umfaßt.
8. Zubereitung nach Anspruch 2, die mikrokristalline Cellulose, trockenfließende Stärke und deren Mischungen als Verdünnungsmittel umfaßt.
9. Zubereitung nach Anspruch 8, die mikrokristalline Cellulose in einer Menge zwischen 24 und 46 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
10. Zubereitung nach Anspruch 9, worin die mikrokristalline Cellulose eine mittlere Teilchengröße von ungefähr 90 Mikron aufweist.
11. Zubereitung nach Anspruch 8, die trockenfließende Stärke in einer Menge von ungefähr 39 Gew.-% bezogen auf

das Gesamtgewicht der Zubereitung umfaßt.

- 5 12. Zubereitung nach Anspruch 2, die Stearylatriumfumarat als Gleitmittel in einer Menge von weniger als 2 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
13. Zubereitung nach Anspruch 2, die kolloidales Siliciumdioxid als Antihafmittel in einer Menge von weniger als 5 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 10 14. Zubereitung nach Anspruch 2, die Aspartam, Natriumsaccharinat oder deren Mischungen als Süßstoff in einer Menge von weniger als 1 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 15 15. Zubereitung nach Anspruch 14, die Natriumsaccharinat in einer Menge zwischen 0,1 und 0,4 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 16 16. Zubereitung nach Anspruch 14, die Aspartam in einer Menge zwischen 0,2 und 0,7 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 17 17. Zubereitung nach Anspruch 2, die Erdbeeraroma in einer Menge zwischen 3 und 5 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 20 18. Zubereitung nach Anspruch 2, die wasserfreie Zitronensäure als Aromastoff in einer Menge zwischen 2 und 4 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 25 19. Zubereitung nach Anspruch 2, die Titandioxid als Trübungsmittel in einer Menge von weniger als 2 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 30 20. Zubereitung nach Anspruch 2, die Rot F, D und C Nr. 3 - Erythrosin als Farbstoff in einer Menge von weniger als 1 Gew.-% bezogen auf das Gesamtgewicht der Zubereitung umfaßt.
- 31 21. Zubereitung nach Anspruch 1, die ein Paar von Verbindungen einschließt, die eine schäumende Wirkung erzeugen.
- 32 22. Zubereitung nach Anspruch 21, worin das die schäumende Wirkung erzeugende Paar aus Zitronensäure und Calciumcarbonat besteht.
- 35

Revendications

- 40 1. Formulation pharmaceutique appropriée pour la préparation de comprimés dispersibles, par compression directe, avec du céfalclore à titre d'ingrédient actif en une quantité entre 35 % et 50 % en poids du poids total de la formulation, avec des excipients et des co-adjuvants appropriés.
- 45 2. Formulation selon la revendication 1, dans laquelle lesdits excipients et co-adjuvants appropriés comprennent les agents désintégrants, les diluants, les lubrifiants, les agents anti-adhérents, les édulcorants, les parfums et facultativement les arômes, les agents opacifiants et les colorants.
- 50 3. Formulation selon la revendication 2, dans laquelle l'agent désintégrant est du glycolate d'amidon de sodium, des dérivés acryliques, des mélanges de glycolate d'amidon de sodium et de carboxyméthylcellulose et, de préférence, de la crospovidone.
- 55 4. Formulation selon la revendication 3, comprenant du glycolate d'amidon de sodium en une quantité entre 5 % et 21 % en poids du poids total de la formulation.
- 56 5. Formulation selon la revendication 3, comprenant des dérivés d'acide acrylique en une quantité d'environ 10% en poids du poids total de la formulation.
- 57 6. Formulation selon la revendication 3, comprenant de la crospovidone en une quantité d'environ 10 % en poids du poids total de la formulation.

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7. Formulation selon la revendication 3, comprenant un mélange de (i) glycolate d'amidon de sodium en une quantité d'environ 14 % en poids du poids total de la formulation, et (ii) carboxyméthylcellulose en une quantité d'environ 10 % en poids du poids total de la formulation, à titre d'agent désintégrant.
- 5 8. Formulation selon la revendication 2, comprenant de la cellulose microcristalline, de l'amidon fluide sec et leurs mélanges, à titre de diluant.
9. Formulation selon la revendication 8, comprenant de la cellulose microcristalline en une quantité entre 24 % et 46 % en poids du poids total de la formulation.
- 10 10. Formulation selon la revendication 9, dans laquelle ladite cellulose microcristalline présente une granulométrie moyenne d'environ 90 µm.
11. Formulation selon la revendication 8, comprenant de l'amidon fluide sec en une quantité d'environ 39 % en poids du poids total de la formulation.
- 15 12. Formulation selon la revendication 2, comprenant du fumarate stéarylique de sodium à titre de lubrifiant, en une quantité inférieure à 2 % en poids du poids total de la formulation.
- 20 13. Formulation selon la revendication 2, comprenant du dioxyde de silicium colloïdal à titre d'agent anti-adhérent, en une quantité inférieure à 5 % en poids du poids total de la formulation.
14. Formulation selon la revendication 2, comprenant de l'aspartame, de la saccharine de sodium ou leurs mélanges à titre d'édulcorant, en une quantité inférieure à 1 % en poids du poids total de la formulation.
- 25 15. Formulation selon la revendication 14, comprenant de la saccharine de sodium, en une quantité entre 0,1 % et 0,4 % en poids du poids total de la formulation.
16. Formulation selon la revendication 14, comprenant de l'aspartame, en une quantité entre 0,2 % et 0,7 % en poids du poids total de la formulation.
- 30 17. Formulation selon la revendication 2, comprenant un parfum de fraise, en une quantité entre 3 % et 5 % en poids du poids total de la formulation.
- 35 18. Formulation selon la revendication 2, comprenant de l'acide citrique anhydre à titre d'arôme, en une quantité entre 2 % et 4 % en poids du poids total de la formulation.
19. Formulation selon la revendication 2, comprenant du dioxyde de titane à titre d'agent opacifiant, en une quantité inférieure à 2 % en poids du poids total de la formulation.
- 40 20. Formulation selon la revendication 2, comprenant de l'Érythrosine - No. 3 Red F, D et C à titre de colorant, en une quantité inférieure à 1 % en poids du poids total de la formulation.
21. Formulation selon la revendication 1, qui comprend une paire de composés produisant un effet effervescent.
- 45 22. Formulation selon la revendication 21, dans laquelle ladite paire effervescente est constituée d'acide citrique et de carbonate de calcium.